

Flexoelectric effect in biaxial nematics

A. Kapanowski

*Institute of Physics, Jagiellonian University,
ulica Reymonta 4, 30-059 Cracow, Poland*

January 12, 2013

Abstract

The flexoelectric (FE) effect provides a linear coupling between electric polarization and orientational deformation in liquid crystals. It influences many electrooptical phenomena and it is used in some bistable nematic devices. A statistical theory of dipole FE polarization in biaxial nematic liquid crystals is used to calculate temperature dependence of order parameters, elastic constants, and FE coefficients. The splitting of the two Meyer FE coefficients and the appearance of new FE coefficients is obtained at the uniaxial-biaxial nematic transition. The ordering of the splitted FE coefficients corresponds to the ordering of the splitted elastic constants.

Keywords: biaxial nematic liquid crystals, flexoelectric effect, elastic constants.

1 Introduction

The biaxial nematic liquid crystals were first predicted by Freiser [1], who showed that molecules with shapes that deviate from cylindrical symmetry could possess a nematic phase with three distinct optical axes. A biaxial nematic phase was first observed in a lyotropic mixture by Yu and Saupe [2] in 1980 but the existence of a thermotropic biaxial system was not certain for many years. Several reports appeared in 2004 on thermotropic liquid crystals formed by side-chain polymers [3], bent-core molecules [4, 5], and tetrapodes [6]. The number of new biaxial systems is constantly growing.

Many theoretical papers [7] and computer simulations [8] show that the molecular shape and pair interaction biaxiality are important for the biaxial phase to exist. However, very often real systems favour packing in the smectic or crystalline biaxial phases. It is a challenge for the theory to find factors responsible for absolute stability of the biaxial nematic phase. It was shown that fluctuations in molecular shape can influence the biaxial nematic phase stability [9]. The motivation for this search ranges from purely academic interest to the potential usage of these materials in faster displays, where in principle the commutation of the secondary director should give lower response times compared to the conventional twisted nematic and ferroelectric smectic devices.

A static electric field imposed on a nematic liquid crystal have many physical effects, but the most important are two of them. One is connected with the anisotropy of the dielectric constant. The second effect is the appearance of the spontaneous polarization in a deformed liquid crystal; this is called the flexoelectric effect. Conversely, an electric field may induce distortions in the bulk. In 1969 Meyer showed that it is a steric effect due to the shape asymmetry of polar molecules [10]. In case of nonpolar molecules the FE effect originates from a gradient of quadrupole moment density [11].

The two FE coefficients were introduced by Meyer for splay and bend distortions of the uniaxial nematic phase [10]. Recently, a statistical theory for the dipole FE polarization was derived in the case of the biaxial nematic phase composed of the C_{2v} molecules [12]. There are six splay-bend deformations of the biaxial nematic phase and thus six FE coefficients are defined, but only five of them are independent. General microscopic expressions for the FE coefficients involve the one-particle distribution function and the potential energy of two-body short-ranged interactions.

The FE effect has a large influence on many phenomena in liquid crystals: electrooptical phenomena and defect formation, for instance. It plays a key role in some device applications. Flexoelectric switching is important in bistable displays [13]-[15]. Flexoelectric coupling in chiral and twisted nematic crystals [16] leads to a linear rotation of the optic axis and also leads to device applications [17]. Flexoelectric coupling in smectic liquid crystals has been shown to stabilize helical structures [18].

The purpose of this study is to calculate the values of the FE coefficients and other material parameters for model systems. The proper form of the interaction potential energy allows us to calculate the temperature dependence of the order parameters, the FE coefficients, and the elastic constants.

The uniaxial and biaxial nematic phases are considered.

2 Description of the system

Let us consider a set of N molecules contained in a volume V , at the temperature T . The molecules are rigid blocks (C_{2v} symmetry) with three translational and three rotational degrees of freedom. It is assumed that the molecules interact via two-body short-range forces that depend on the distance between the molecules ($\vec{u} = \vec{r}_2 - \vec{r}_1 = u\vec{\Delta}$) and their orientations described by the three Euler angles $R = (\phi, \theta, \psi)$ or three orthonormal vectors $(\vec{l}, \vec{m}, \vec{n})$.

The microscopic free energy F for the system is given by

$$F = F_{ent} + F_{int}, \quad (1)$$

$$\beta F_{ent} = \int d\vec{r} dR G(\vec{r}, R) \{\ln[G(\vec{r}, R)\Lambda] - 1\}, \quad (2)$$

$$\beta F_{int} = -\frac{1}{2} \int d\vec{r}_1 dR_1 d\vec{r}_2 dR_2 G(\vec{r}_1, R_1) G(\vec{r}_2, R_2) f_{12}. \quad (3)$$

Here $f_{12} = \exp(-\beta\Phi_{12}) - 1$ is the Mayer function, Φ_{12} the potential energy of interactions, $dR = d\phi d\theta \sin\theta d\psi$, $\beta = 1/(k_B T)$, and Λ is related to the ideal gas properties. The one-particle distribution function G has the normalization

$$\int d\vec{r} dR G(\vec{r}, R) = N. \quad (4)$$

The equilibrium distribution G minimizing the free energy (1) satisfies the equation

$$\ln[G(\vec{r}_1, R_1)\Lambda] - \int d\vec{r}_2 dR_2 G(\vec{r}_2, R_2) f_{12} = \text{const}. \quad (5)$$

For the homogeneous phase $G = G_0$ does not depend on the position and it has the form [19]

$$G_0(R) = G_0(\vec{l} \cdot \vec{L}, \vec{l} \cdot \vec{N}, \vec{n} \cdot \vec{L}, \vec{n} \cdot \vec{N}), \quad (6)$$

where the orthonormal vectors $(\vec{L}, \vec{M}, \vec{N})$ define the biaxial nematic phase axes. In practice we characterize the alignment not through the full function G , but by some numerical parameters - order parameters. In the case of

the biaxial nematic phase the main order parameters are the orientational distribution averages of the following four functions [20]: $F_{00}^{(2)}$, $F_{02}^{(2)}$, $F_{20}^{(2)}$, and $F_{22}^{(2)}$. We note that there are other notations [22]. In the uniaxial nematic phase the functions $F_{00}^{(2)}$ and $F_{02}^{(2)}$ have nonzero averages only. The molecule alignment can be also described by order tensors Q which are often calculated for computer simulations [23] where the molecular and laboratory axes must be distinguished. The order tensors are defined as

$$Q_{\alpha\beta}^{ll} = (3l_\alpha l_\beta - \delta_{\alpha\beta})/2, \quad (7)$$

$$Q_{\alpha\beta}^{mm} = (3m_\alpha m_\beta - \delta_{\alpha\beta})/2, \quad (8)$$

$$Q_{\alpha\beta}^{nn} = (3n_\alpha n_\beta - \delta_{\alpha\beta})/2. \quad (9)$$

3 Elastic deformations of the phase

Oriental ordering of biaxial nematics is usually described by the three orthonormal vectors

$$\vec{L} = R_{1\alpha}\vec{e}_\alpha, \vec{M} = R_{2\alpha}\vec{e}_\alpha, \vec{N} = R_{3\alpha}\vec{e}_\alpha. \quad (10)$$

In the homogeneous phase the vectors $(\vec{L}, \vec{M}, \vec{N})$ are constant in space, but in a deformed phase they depend on the position in space. In a continuum approach the distortion free-energy density f_d is obtained as an expansion about an undistorted reference state with respect to gradients of the vectors $(\vec{L}, \vec{M}, \vec{N})$. The form of the f_d can be derived in many alternative ways but we use the form presented by Stallinga and Vertogen [21] (the surface terms are neglected)

$$\begin{aligned} f_d = & \frac{1}{2}K_{1111}(D_{11})^2 + \frac{1}{2}K_{1212}(D_{12})^2 + \frac{1}{2}K_{1313}(D_{13})^2 \\ & + \frac{1}{2}K_{2121}(D_{21})^2 + \frac{1}{2}K_{2222}(D_{22})^2 + \frac{1}{2}K_{2323}(D_{23})^2 \\ & + \frac{1}{2}K_{3131}(D_{31})^2 + \frac{1}{2}K_{3232}(D_{32})^2 + \frac{1}{2}K_{3333}(D_{33})^2 \\ & + K_{1122}D_{11}D_{22} + K_{1133}D_{11}D_{33} + K_{2233}D_{22}D_{33} \\ & + K_{1221}D_{12}D_{21} + K_{1331}D_{13}D_{31} + K_{2332}D_{23}D_{32}. \end{aligned} \quad (11)$$

$$D_{ij} = \frac{1}{2}\epsilon_{jkl}R_{i\alpha}R_{k\beta}\partial_\alpha R_{l\beta}. \quad (12)$$

Microscopic expressions for the elastic constants K_{ijkl} were derived in [19] and it was shown that there are 12 independent bulk constants because

$$K_{1221} = K_{1122}, \quad K_{1331} = K_{1133}, \quad K_{2332} = K_{2233}. \quad (13)$$

4 Flexoelectric polarization

Liquid crystalline phases often consist of polar molecules but in homogeneous nematic phases the average polarization is zero. On the other hand, a phase distortion can produce a polarization and this is called the FE effect. In a continuum approach the FE polarization of the biaxial nematic phase depends on the spatial derivatives of the vectors $(\vec{L}, \vec{M}, \vec{N})$ [12]

$$P_\alpha = \sum_i (s_{ii} R_{i\alpha} \partial_\beta R_{i\beta} + b_{ii} R_{i\beta} \partial_\beta R_{i\alpha}). \quad (14)$$

The parameters s_{ii} and b_{ii} , ($i = 1, 2, 3$) are not unique because if we add any constant to all of them, the polarization will not change. The physical FE coefficients a_i ($i = 4, \dots, 9$) are

$$\begin{aligned} a_4 &= s_{33} - b_{11}, & a_5 &= s_{22} - b_{11}, & a_6 &= s_{33} - b_{22}, \\ a_7 &= s_{11} - b_{22}, & a_8 &= s_{22} - b_{33}, & a_9 &= s_{11} - b_{33}. \end{aligned} \quad (15)$$

The coefficients satisfy the identity

$$a_4 - a_5 - a_6 + a_7 + a_8 - a_9 = 0. \quad (16)$$

Deformations of the biaxial nematic phase connected with the FE effect are given in Table 1. In the case of the uniaxial nematic phase the FE polarization has the form

$$P_\alpha = e_1 N_\alpha \partial_\beta N_\beta + e_3 N_\beta \partial_\beta N_\alpha. \quad (17)$$

Let us define the molecule electric dipole moment as

$$\mu_\alpha = \mu_1 l_\alpha + \mu_2 m_\alpha + \mu_3 n_\alpha. \quad (18)$$

In the case of the molecular interactions described below, the FE coefficients can be expressed as follows

$$a_4 = \int d\vec{u} dR_1 dR_2 f_{12} G_0(R_1) \mu_3 n_{1z} (-u_x) (U_{2z} - W_{2x}), \quad (19)$$

$$a_5 = \int d\vec{u} dR_1 dR_2 f_{12} G_0(R_1) \mu_3 n_{1y} (-u_x) U_{2y}, \quad (20)$$

$$a_6 = \int d\vec{u} dR_1 dR_2 f_{12} G_0(R_1) \mu_3 n_{1z} u_y W_{2y}, \quad (21)$$

$$a_7 = \int d\vec{u} dR_1 dR_2 f_{12} G_0(R_1) \mu_3 n_{1x} u_y U_{2y}, \quad (22)$$

$$a_8 = \int d\vec{u} dR_1 dR_2 f_{12} G_0(R_1) \mu_3 n_{1y} (-u_z) W_{2y}, \quad (23)$$

$$a_9 = \int d\vec{u} dR_1 dR_2 f_{12} G_0(R_1) \mu_3 n_{1x} u_z (U_{2z} - W_{2x}), \quad (24)$$

where it is assumed that \vec{n} defines the molecule C_2 axis and

$$U_\alpha = \partial_1 G_0 l_\alpha + \partial_3 G_0 n_\alpha, \quad W_\alpha = \partial_2 G_0 l_\alpha + \partial_4 G_0 n_\alpha. \quad (25)$$

5 Results

We performed our calculations for the square-well potential energy of the form

$$\Phi_{12}(u/\sigma) = \begin{cases} +\infty & \text{for } (u/\sigma) < 1, \\ -\epsilon & \text{for } 1 < (u/\sigma) < 2, \\ 0 & \text{for } (u/\sigma) > 2, \end{cases} \quad (26)$$

where ϵ is the depth of the well and σ depends on the molecule orientations and on the vector $\vec{\Delta}$

$$\begin{aligned} \sigma = & \sigma_0 + \sigma_1 (\vec{\Delta} \cdot \vec{n}_1 - \vec{\Delta} \cdot \vec{n}_2) + \sigma_2 [(\vec{\Delta} \cdot \vec{n}_1)^2 + (\vec{\Delta} \cdot \vec{n}_2)^2] \\ & + \sigma_3 [(\vec{\Delta} \cdot \vec{l}_1)^2 + (\vec{\Delta} \cdot \vec{l}_2)^2] \end{aligned} \quad (27)$$

The parameter σ_0 defines the length scale, σ_1 defines the FE term, σ_2 and σ_3 define biaxial nematic terms. We used the density $NV_{mol}/V = 0.1$, the molecule volume V_{mol} was estimated from the mutually excluded volume. The FE coefficients were expressed in μ_i/σ_0^2 , the elastic constants in ϵ/σ_0 , and the temperature in ϵ/k_B . The parameters σ_i are given in Table 2. The two physical systems are considered that consist of wedge-shaped and banana-shaped molecules.

5.1 Wedge-shaped biaxial molecules

In the system of wedge-shaped biaxial molecules, on decreasing the temperature we meet the first order transition from the isotropic to the uniaxial

nematic phase at $T_{IN} = 0.618$ and the second order transition to the biaxial nematic phase at $T_{NB} = 0.401$. The temperature dependence of the order tensors is presented in Fig. 1. The temperature dependence of the elastic constants and the FE coefficients are presented in Figs. 2 and 3, respectively.

The values of Q_{zz}^{nn} show that long molecule axes are directed along the Z axis in the whole nematic region, whereas the values of Q_{xx}^{ll} reveal the alignment of short molecule axes along the X axis and it is enhanced in the biaxial nematic phase. The splay elastic constant K_1 splits into K_{1212} and K_{2121} . The bend elastic constant K_3 splits into K_{3232} and K_{3131} . Note that the equality $K_1 = K_3$ is accidental and results from neglecting order parameters $F_{\mu\nu}^{(j)}$ with $j > 2$. The splay FE coefficient e_1 splits into a_4 and a_6 ($a_4 > a_6 > 0$). The bend FE coefficient $-e_3$ splits into a_8 and a_9 ($0 > a_8 > a_9$). The coefficients a_5 and a_7 are small and almost always negative.

5.2 Banana-shaped biaxial molecules

In the system of banana-shaped biaxial molecules, on decreasing the temperature we meet the first order transition from the isotropic to the uniaxial nematic phase at $T_{IN} = 0.595$ and the second order transition to the biaxial nematic phase at $T_{NB} = 0.382$. The temperature dependence of the order tensors is presented in Fig. 4. The temperature dependence of the elastic constants and the FE coefficients are presented in Figs. 5 and 6, respectively.

According to the values of Q_{zz}^{ll} long molecule axes are directed along the Z axis in the whole nematic region, whereas the values of Q_{xx}^{nn} show the alignment of short molecule axes along the X axis and it is enhanced in the biaxial nematic phase. The behaviour of the elastic constants is similar to the case of the wedge-shaped molecules because the FE term is small in both cases. The bend FE coefficient $-e_3$ splits into a_8 and a_9 ($0 > a_8 > a_9$). The splay FE coefficient e_1 is smaller than e_3 and it splits into a_4 and a_6 . The coefficients a_5 and a_7 are again small but comparable with a_4 and a_6 . The sign of some coefficient can change on changing the temperature.

6 Conclusions

In this paper, the statistical theory was used to study the temperature dependence of the order parameters, elastic constants, and FE coefficients of biaxial nematic liquid crystals. In order to calculate these macroscopic pa-

rameters one needs the one-particle distribution function and the potential energy of molecular interactions. The two physical systems were considered. The splittings of the FE coefficients and the elastic constants were obtained at the uniaxial-biaxial nematic transition. New small FE coefficients appeared at the transition. The ordering of the splitted FE coefficients corresponds to the ordering of the splitted elastic constants.

The FE coefficients were proportional to the dipole moment component parallel to the molecule C_{2v} symmetry axis. This was the result of the interactions potential symmetry. The behaviour of the main FE coefficients, e_1 for the wedge-shaped molecules and e_3 for the banana-shaped molecules, is clear and it is in the agreement with previous studies [24]. On the other hand, it seems that other FE coefficients should be interpreted with caution. It is possible that higher order parameters can have a significant contribution.

At present stage, the direct comparison between the theory and the experiment is not possible because to our knowledge the FE coefficients have not been measured for the biaxial nematic phase. What is more, even for the uniaxial nematic phase the experimental data are still scarce and sometimes contradictory [25]. However, when biaxial nematic phases become more widespread, the presented theory will be helpful in practical applications.

Acknowledgements

The author is grateful to J. Spálek for his support and discussions.

References

- [1] M. J. Freiser, "Ordered states of a nematic liquid", *Phys. Rev. Lett.* **24**, 1041-1043 (1970).
- [2] L. J. Yu and A. Saupe, "Observation of a biaxial nematic phase in potassium-1-decanol-water mixtures", *Phys. Rev. Lett.* **45**, 1000-1003 (1980).
- [3] K. Severing and K. Saalwachter, "Biaxial nematic phase in a thermotropic liquid crystalline side-chain polymer", *Phys. Rev. Lett.* **92**, 125501 (2004).

- [4] L. A. Madsen, T. J. Dingemans, M. Nakata, and E. T. Samulski, "Thermotropic biaxial nematic liquid crystals", *Phys. Rev. Lett.* **92**, 145505 (2004).
- [5] B. R. Acharya, A. Primak, and S. Kumar, "Biaxial nematic phase in bent-core thermotropic mesogens", *Phys. Rev. Lett.* **92**, 145506 (2004).
- [6] K. Merkel, A. Kocot, J. K. Vij, R. Korlacki, G. H. Mehl, and T. Meyer, "Thermotropic biaxial nematic phase in liquid crystalline organo-siloxane tetrapodes", *Phys. Rev. Lett.* **93**, 237801 (2004).
- [7] L. Longa, P. Grzybowski, S. Romano, and E. Virga, "Minimal coupling model of the biaxial nematic phase", *Phys. Rev.* **E 71**, 051714 (2005).
- [8] R. Berardi, L. Muccioli, S. Orlandi, M. Ricci, and C. Zannoni, "Computer simulations of biaxial nematics", *J. Phys.: Condens. Matter* **20**, 463101 (2008).
- [9] L. Longa, G. Pająk, and T. Wydro, "Stability of biaxial nematic phase for systems with variable molecular shape anisotropy", *Phys. Rev.* **E 76**, 011703 (2007).
- [10] R. B. Meyer, "Piezoelectric effects in liquid crystals", *Phys. Rev. Lett.* **22**, 918-921 (1969).
- [11] J. Prost and J. P. Marcerou, "On the microscopic interpretation of flexoelectricity", *J. Phys. (Paris)* **38**, 315-324 (1977).
- [12] A. Kapanowski, "Flexoelectric polarization in the biaxial nematic phase", *Phys. Rev.* **E 77**, 052702 (2008).
- [13] C. Denniston and J. M. Yeomans, "Flexoelectric surface switching of bistable nematic devices", *Phys. Rev. Lett.* **87**, 275505 (2001).
- [14] A. J. Davidson and N. J. Mottram, "Flexoelectric switching in a bistable nematic device", *Phys. Rev.* **E 65**, 051710 (2002).
- [15] L. A. Parry-Jones, R. B. Meyer, and S. J. Elston, "Mechanisms of flexoelectric switching in a zenithally bistable nematic device", *J. Appl. Phys.* **106**, 014510 (2009).

- [16] J. S. Patel and R. B. Meyer, "Flexoelectric electro-optics of a cholesteric liquid crystal", *Phys. Rev. Lett.* **58**, 1538-1540 (1987).
- [17] A. E. Blatch, M. J. Coles, B. Musgrave, and H. J. Coles, "Flexoelectric liquid crystal bimesogens", *Mol. Cryst. Liq. Cryst.* **401**, 161-169 (2003).
- [18] M. Cepic and B. Zeks, "Flexoelectricity and Piezoelectricity: The Reason for the Rich Variety of Phases in Antiferroelectric Smectic Liquid Crystals", *Phys. Rev. Lett.* **87**, 085501 (2001).
- [19] A. Kapanowski, "Statistical theory of elastic constants of biaxial nematic liquid crystals", *Phys. Rev.* **E 55**, 7090-7104 (1997).
- [20] M. Fialkowski, A. Kapanowski, K. Sokalski, "Microscopic approach to theory of biaxial nematic liquid crystals", *Mol. Cryst. Liq. Cryst.* **265**, 371-385 (1995).
- [21] S. Stallinga, G. Vertogen, "Theory of orientational elasticity", *Phys. Rev.* **E 49**, 1483-1495 (1994).
- [22] R. Rosso, "Orientational order parameters in biaxial nematics: Polymorphic notation", *Liq. Cryst.* **34**, 737-748 (2007).
- [23] P. J. Camp and M. P. Allen, "Phase diagram of the hard biaxial ellipsoid fluid", *J. Chem. Phys.* **106**, 6681 (1997).
- [24] A. Kapanowski, "Flexoelectric effect modelling", *Opto-Electron. Rev.* **16**, 9-12 (2008).
- [25] A. G. Petrov, Measurements and interpretation of flexoelectricity", in *Physical properties of liquid crystals: nematics*, pp. 251-264, edited by D. Dunmur, A. Fukuda, and G. L. Luckhurst, INSPEC, The Institution of Electrical Engineers, London, 2001.

Table 1: Deformations of the biaxial nematic phase connected with the FE effect. The corresponding elastic constants and the FE coefficients are given, the values for the uniaxial nematic phase are in parentheses.

Deformation	Elastic constant	FE coefficient
\vec{N} splay, \vec{L} bend	K_{1212} (K_1)	a_4 (e_1)
\vec{M} splay, \vec{L} bend	K_{1313} (0)	a_5 (0)
\vec{N} splay, \vec{M} bend	K_{2121} (K_1)	a_6 (e_1)
\vec{L} splay, \vec{M} bend	K_{2323} (0)	a_7 (0)
\vec{M} splay, \vec{N} bend	K_{3131} (K_3)	a_8 ($-e_3$)
\vec{L} splay, \vec{N} bend	K_{3232} (K_3)	a_9 ($-e_3$)

Table 2: Parameters σ_i used in calculations.

Molecules	σ_1/σ_0	σ_2/σ_0	σ_3/σ_0	Long axis	Short axis	C_2 axis
wedge-like	0.2	0.5	-0.4	\vec{n}	\vec{l}	\vec{n}
banana-like	0.2	-0.4	0.5	\vec{l}	\vec{n}	\vec{n}

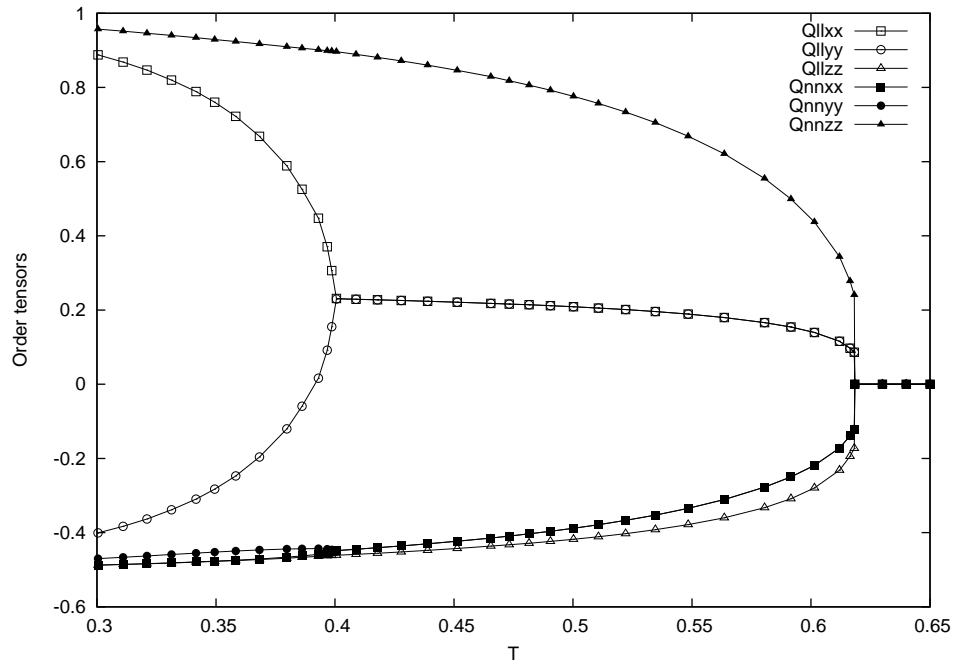


Figure 1: The temperature dependence of the order tensors for wedge-shaped biaxial molecules.

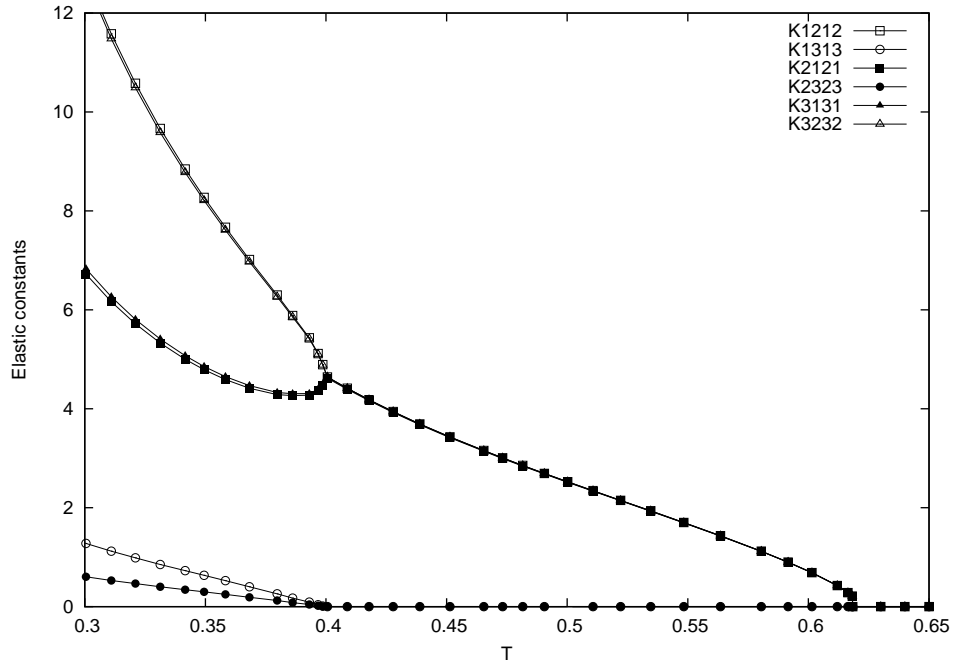


Figure 2: The temperature dependence of the elastic constants for wedge-shaped biaxial molecules. The squares, triangles, and circles denote deformations with \vec{N} splay, \vec{N} bend, and \vec{N} constant, respectively. The empty (filled) symbols indicate the larger (smaller) parameter.

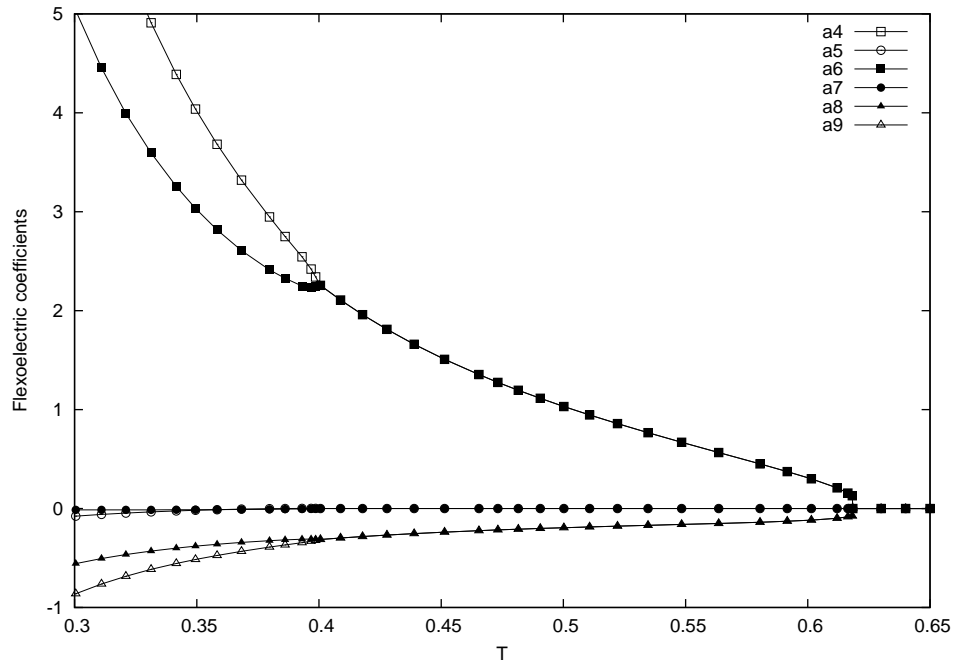


Figure 3: The temperature dependence of the flexoelectric coefficients for wedge-shaped biaxial molecules. Symbols have the same meaning as in Fig. 2.

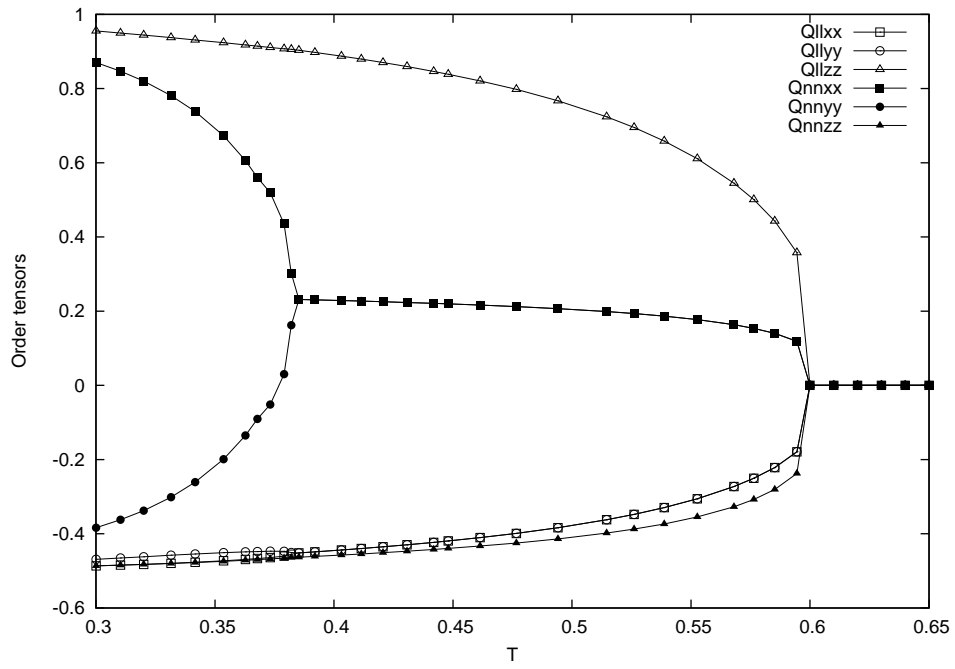


Figure 4: The temperature dependence of the order tensors for banana-shaped biaxial molecules.

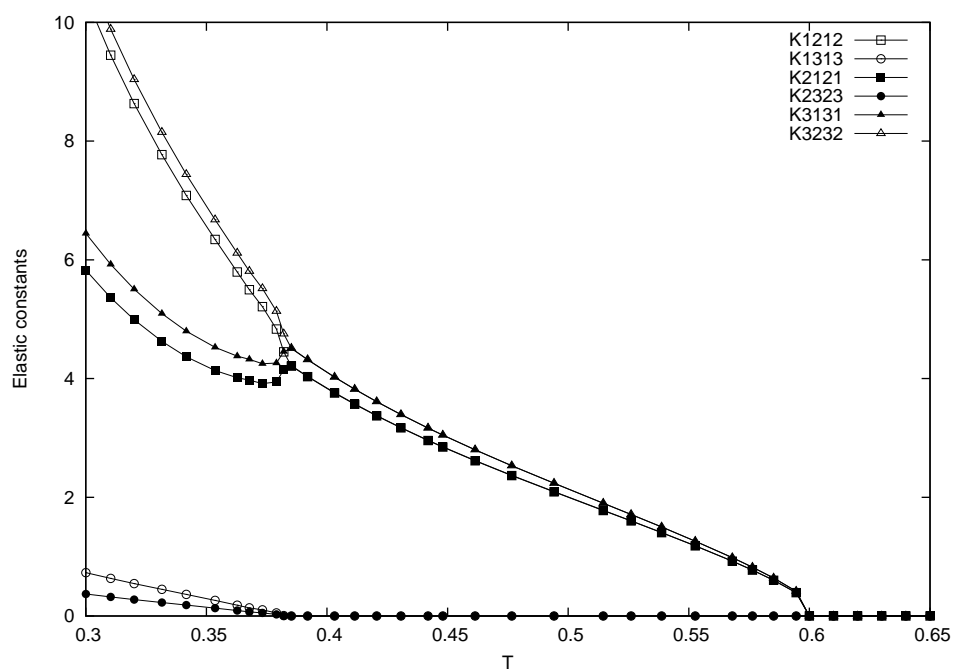


Figure 5: The temperature dependence of the elastic constants for banana-shaped biaxial molecules. Symbols have the same meaning as in Fig. 2.

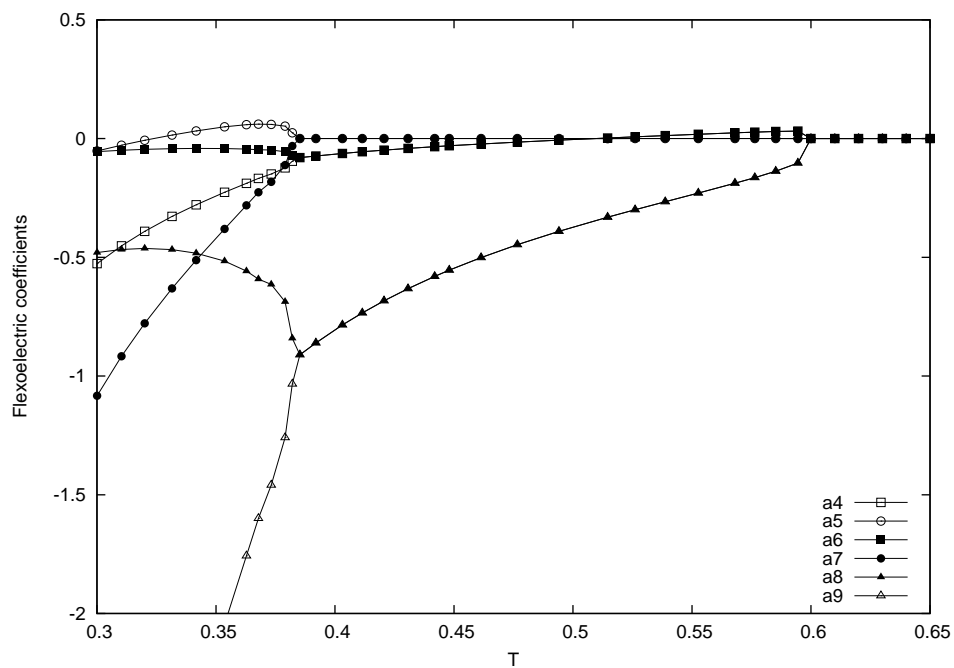


Figure 6: The temperature dependence of the flexoelectric coefficients for banana-shaped biaxial molecules. Symbols have the same meaning as in Fig. 2.